

IMPORTANT NAME CHANGE ANNOUNCEMENT

Larotid (amoxicillin)

is the new name for Larocin

Since its introduction in March of 1974, Larocin has been prescribed more than a million times by physicians in the United States. In several of these instances, written prescriptions for Larocin have been confused with Lanoxin, Burroughs Wellcome Company's brand of digoxin. Although the reported incidence of such confusion has been extremely low, Roche Laboratories has changed the name of its product to LAROTID (amoxicillin). We hope you will agree that this action is in the best interest of the patient and of everyone concerned.

Before prescribing, please consult complete product information, a summary of which follows:

Indications: Infections due to susceptible strains of the following gram-negative organisms: *H. influenzae*, *E. coli*, *P. mirabilis* and *N. gonorrhoeae*; and gram-positive organisms: streptococcal (including *Streptococcus faecalis*), *D. pneumoniae* and non-penicillase-producing staphylococci. Therapy may be instituted prior to obtaining results from bacteriological and susceptibility studies to determine causative organisms and susceptibility to amoxicillin.

Contraindications: In individuals with history of allergic reaction to penicillins.

Warnings: SERIOUS AND OCCASIONALLY FATAL HYPERSENSITIVITY (ANAPHYLACTOID) REACTIONS REPORTED IN PATIENTS ON PENICILLIN THERAPY. ALTHOUGH MORE FREQUENT FOLLOWING PARENTERAL THERAPY, ANAPHYLAXIS HAS OCCURRED IN PATIENTS ON ORAL PENICILLINS. MORE LIKELY IN INDIVIDUALS WITH HISTORY OF SENSITIVITY TO MULTIPLE ALLERGENS. BEFORE THERAPY, INQUIRE CONCERNING PREVIOUS HYPERSENSITIVITY REACTIONS TO PENICILLINS, CEPHALOSPORINS OR OTHER ALLERGENS. IF ALLERGIC REACTION OCCURS, INSTITUTE APPROPRIATE THERAPY AND CONSIDER DISCONTINUANCE OF AMOXICILLIN. SERIOUS ANAPHYLACTOID REACTIONS REQUIRE IMMEDIATE EMERGENCY TREATMENT WITH EPINEPHRINE, ADMINISTER OXYGEN, INTRAVENOUS STEROIDS AND AIRWAY MANAGEMENT, INCLUDING INTUBATION, AS INDICATED.

Usage in Pregnancy: Safety in pregnancy not

established.

Precautions: As with any potent drug, assess renal, hepatic and hematopoietic function periodically during prolonged therapy. Keep in mind possibility of superinfections with mycotic or bacterial pathogens; if they occur, discontinue drug and/or institute appropriate therapy.

Adverse Reactions: As with other penicillins, unoward reactions will likely be essentially limited to sensitivity phenomena and more likely occur in individuals previously demonstrating penicillin hypersensitivity and those with history of allergy, asthma, hay fever or urticaria. Adverse reactions reported as associated with use of penicillins: Gastrointestinal: Nausea, vomiting, diarrhea. Hypersensitivity Reactions: Erythematous maculopapular rashes, urticaria. NOTE: Urticaria, other skin rashes and serum sickness-like reactions may be controlled with antihistamines and, if necessary, systemic corticosteroids. Discontinue amoxicillin unless condition is believed to be life-threatening and amenable only to amoxicillin therapy. Liver: Moderate rise in SGOT noted, but significance unknown. Hematologic and Lymphatic Systems: Anemia, thrombocytopenia, thrombocytopenic purpura, eosinophilia, leukopenia, agranulocytosis. All are usually reversible on discontinuation of therapy and believed to be hypersensitivity phenomena.

Dosage: Ear, nose, throat, cutaneous tract, skin and soft tissue infections—Adults: 250 mg every 8 hours. Children: 20 mg/kg/day in divided doses every 8 hours; under 8 kg, 0.5 ml of Pediatric Drops every 8 hours; 8-6 kg, 1 ml of Pediatric Drops every 8 hours. Lower respiratory tract infections and se-

vere infections or those caused by less susceptible organisms—Adults: 500 mg every 8 hours. Children: 40 mg/kg/day in divided doses every 8 hours; under 8 kg, 1 ml of Pediatric Drops every 8 hours; 8-6 kg, 2 ml of Pediatric Drops every 8 hours. Gonorrhea (acute uncomplicated nongonococcal and urethral infections)—Males and females: 3 grams as a single oral dose. NOTE: Children weighing more than 8 kg should receive appropriate dose of oral suspension 125 mg or 250 mg/5 ml. Children weighing 20 kg or more should be dosed according to adult recommendations.

Notes: In gonorrhea with suspected lesion of syphilis, perform dark-field examinations before amoxicillin therapy and monthly serological tests for at least four months. In chronic urinary tract infections, frequent bacteriological and clinical appraisals are necessary. Smaller than recommended doses should not be used. In stubborn infections, several weeks' therapy may be required. Except for gonorrhea, continue treatment for a minimum of 48-72 hours after patient is asymptomatic or bacterial eradication is evidenced. Treat hemolytic streptococcal infections for at least 10 days to prevent acute rheumatic fever or glomerulonephritis.

Supplied: Amoxicillin as the trihydrate: Capsules, 250 mg and 500 mg; oral suspension, 125 mg/5 ml and 250 mg/5 ml; pediatric drops, 50 mg/ml.

Roche Laboratories
Division of Hoffmann-La Roche Inc.
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Med Trib 39

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world news of medicine and its practice—fast, accurate, complete

and Medical News—
Wednesday, November 5, 1975

From Planned Parenthood:

New Guidelines Set For Contraception In Women Over 40

By FRANCES GOODNIGHT
Medical Tribune Staff

DETROIT—The first successful total correction of transposed great arteries was reported here by a Brazilian surgical team.

Overcoming technical problems that have frustrated heart surgeons for more than two decades, the team was able to transfer the position of the coronary

arteries in retransposed great vessels and achieve normal blood flow conduits in a 40-day-old infant with a large ventricular septal defect, Dr. A. D. Jatene told the 2nd International Symposium on Cardiac Surgery at the Henry Ford Hospital.

Dr. Jatene is Professor of Surgery at the Cardiologic Institute in Sao Paulo, Brazil.

The achievement was described as a "great technical triumph" by Dr. John W. Kirklin, Professor and Chairman of the Department of Surgery, University of Alabama. He added that Dr. Jatene's procedure offers "a very exciting" surgical approach, especially in patients with a large VSD.

In describing the new procedure, Dr.

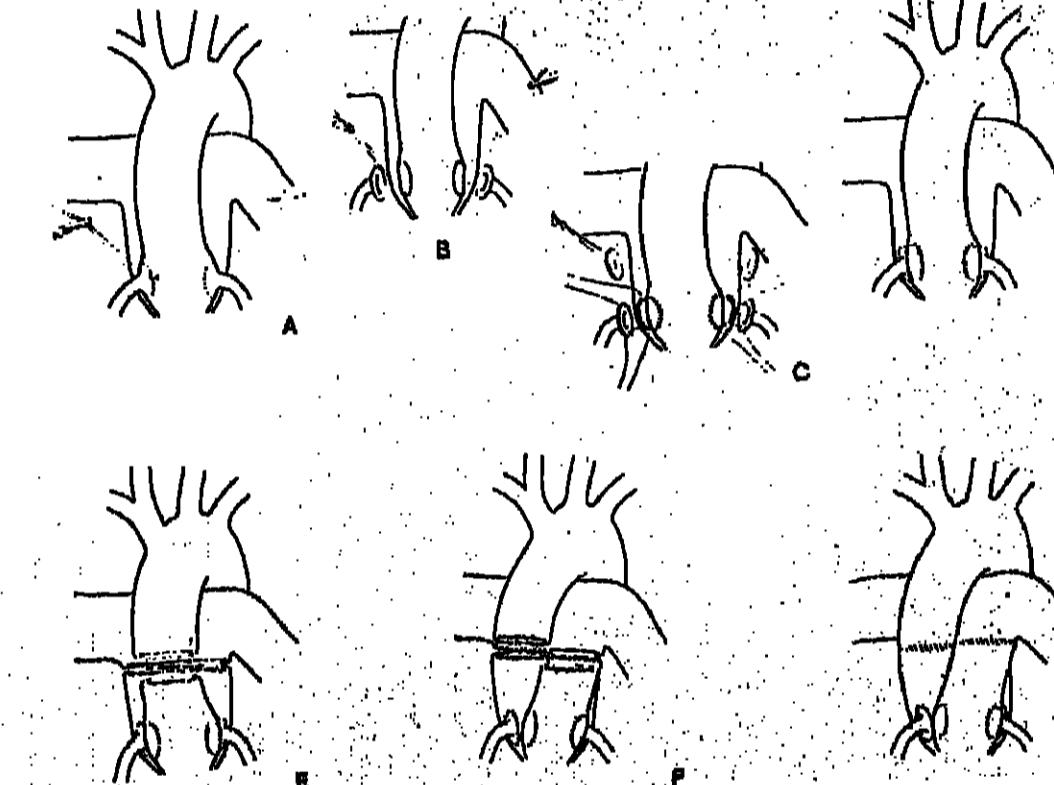
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Transposed Arteries: First Total Correction

By NATHAN HORWITZ
Medical Tribune Staff

DETROIT—The first successful total correction of transposed great arteries was reported here by a Brazilian surgical team.

Overcoming technical problems that have frustrated heart surgeons for more than two decades, the team was able to transfer the position of the coronary



Schematic presentation of the new procedure for total anatomical correction of transposed great vessels in patients with VSD. Figure (A) shows ascending aorta and pulmonary trunk are resected (E), with differences in diameter between the vessels corrected by two sutures in the distal and proximal stumps of pulmonary artery. Distal end of pulmonary artery is sutured to proximal end of anterior artery, now without coronary arteries.

Major Victory Seen In Capitation Grant Policy Reversal

High Cataract Rate Found in Child Asthmatics on Steroids

Medical Tribune Report

DENVER—A high incidence of cataract formation in asthmatic children who regularly take corticosteroids was reported here by a team of physicians from Fitzsimons Army Medical Center and the National Asthma Center.

The study, presented at the 28th Annual Symposium on Pulmonary Diseases here, revealed cataract formation in 10.8 per cent of 92 long-standing severe, steroid-dependent asthmatics.

The policy change is viewed by observers as a major victory by new H.E.W. Secretary F. David Mathews, Ph.D., and Dr. Theodore Cooper, Assistant Secretary for Health, over the Office of Management and Budget.

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to the association of chronic steroid use in child asthmatics with a change in lens pathology.

"Overall, the children in our study ranged from 11 to 15 years of age with an average five-year history of steroid dependency. Slitlamp examination by two independent ophthalmologists revealed definite cataract formation in 10 children, with an additional 21 patients showing some change in lens pathology.

"Obviously, some of these children wouldn't be alive or couldn't function normally without steroids, despite the advances made in chemotherapeutic agents. However, we think physicians should be made aware of our findings," Dr. Spaulding told MEDICAL TRIBUNE.

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New Drug Combination Held Effective Against All Bacteria

Medical Tribune Report

WASHINGTON, D.C.—Development of a new antibacterial combination drug which has proven effective against every bacterial species tested so far, including *Pseudomonas aeruginosa*, was reported here by investigators from the Merck Sharp & Dohme Research Laboratories at a conference sponsored by the American Society for Microbiology.

While the drug has not as yet been studied in man, extensive laboratory and animal tests suggest it may be a potential alternative agent in the treatment of bacterial strains that have acquired a resistance to other antibiotics, the scientists said.

Guarded Reaction

Initial reaction to the report was guarded. Reflecting the opinion of many, Dr. Merrill Snyder, Professor of Medicine in Clinical Microbiology at the University of Maryland School of Medicine in Baltimore, said, "While I commend the investigators on their work, the results are far from applicable to man. The concepts that have been presented are certainly intriguing but whether this will have some practical application remains to be seen."

MK641/MK642, as the drug is currently known, works by inhibiting bacterial cell wall biosynthesis, explained Frederick M. Kahan, discoverer of the

antibacterial combination.

Although bacterial cell wall biosynthesis is also the target of attack of several classes of widely-used antibiotics, including the penicillins and cephalosporins, the new agent is chemically unrelated. It is a fixed ratio combination of 2-deutero-3-fluoro-D-alanine (DFA), a new substance synthesized by Merck scientists, and a derivative of cycloserine (PCS), a 20-year-old antibiotic with limited therapeutic applications.

When combined, the two elements work synergistically to prevent bacteria from synthesizing D-alanine, an indispensable constituent of the cell wall of every type of bacteria, the research team explained.

Development of MK641/MK642 derived from observations that bacteria produce D-alanine through enzymatic conversion of L-alanine and that while L-alanine plays a key role in human metabolism, D-alanine does not. Therefore, the team theorized, an agent which prevented production of only D-alanine would eliminate bacteria in an infected individual without interfering with normal body functioning.

Biologic analyses of DFA show that it prevents bacteria from synthesizing the necessary D-alanine. However, DFA in concentrations several times higher than the minimum inhibitory



Japan's Empress Nagako (right rear) looks on as patients at Chicago's Wyler Children's Hospital prepare for surgery.

concentration has the paradoxical ability to restore bacteria to normal growth. In a phenomenon called "self-reversal," DFA is used by bacteria in place of the missing D-alanine, Mr. Kahan told the meeting.

The addition of PCS, he continued, successfully prevents bacteria from using DFA and thereby ensures DFA's

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Careful Drug Use Urged in Intractable Pain

Medical Tribune World Service

FLORENCE, ITALY—The use of adjuvant drugs such as anti-anxiety and ataractic agents in tandem with narcotics when the latter become necessary for relief of intractable pain was recommended here by Dr. Francis F. Foldes, Professor of Anesthesiology at the Albert Einstein College of Medicine.

"By the judicious combination of these agents, it is possible to provide active, pain-free, and relatively alert days and restful nights for the patient," Dr. Foldes told the First World Congress on Pain Research and Therapy.

Dr. Foldes cautioned against administration of narcotics for painful conditions of limited duration. Pointing out that "even relatively brief" use may cause physical and psychologic dependence, he advises that narcotic use in the presence of acute pain be limited to emergency situations (extensive trauma, acute coronary occlusion, etc.) and severe postoperative pain.

The anesthesiologist also emphasized the need to avoid narcotics for as long as possible in patients with chronic pain and relatively long life expectancy.

"All other methods of pain relief such as the use of analgesics, chemical or surgical interruption of pain pathways, self-induced electrical stimulation of the spinal cord or specific brain areas, and various forms of psychotherapy should be tried before resorting to the chronic administration of narcotics," he said.

But when intractable pain makes the use of narcotics unavoidable, Dr.

Foldes suggests the following guidelines for management of patients who can remain at home:

Management Guides

- Start with orally active narcotics—specifically, the less potent compounds like codeine or oxycodone—and increase dosage gradually. With some types of pain, the pain reduction can be potentiated by non-narcotic analgesics.
- If untoward side effects develop, try first an orally active agonist-antagonist (such as pentazocine) and subsequently meperidine, methadone, or levorphan.
- Try to increase the analgesic effect of narcotics, and diminish their side effects, by giving adjuvant drugs. For example, during the daytime, benefits may be gained by the simultaneous administration of dextroamphetamine sulfate with the narcotic. At night, the combination of the narcotic with a suitable ataractic agent will produce analgesia and sleep "with less respiratory depression than that encountered after larger doses of narcotics used alone."
- In the presence of fear, anxiety, and depression, employ a combination of narcotics with a suitable psychotropic agent. Tranquillizers often relieve narcotic-induced nausea and vomiting, although they are less effective against such symptoms caused by the pathologic process itself.
- Consider anticholinesterases to combat constipation and urinary retention, since they permit reduction of the narcotic dosage and have a direct

stimulating effect on gastrointestinal motility and on bladder tone.

• Keep in mind the possibility that adjuvant drugs can produce untoward effects—either on their own or by interaction with narcotics. "The most important of these are the orthostatic hypotension caused by many ataractic drugs and the markedly increased respiratory depression resulting from the combined use of monoamine oxidase inhibitors and narcotics."

Absolutely Wrong!

The final instruction is to follow the juice with butter or a cooking oil which he calls "absolutely wrong," since it will inhibit the physician's ability to determine the extent of injury and what further treatment is necessary."

Another instance of an outdated antidote directive, according to Dr. Rumack, is the caution found on the labels of most petroleum distillates, oils, and other hydrocarbons, to avoid induction of vomiting.

He said that most major poison centers now follow the policy of inducing emesis if the patient has ingested a possibly toxic amount of hydrocarbon. Management of pain in this setting depends on the circumstances, he pointed out. For patients who are not bedridden, he believes the principles governing the use of narcotics should be similar to those advised for care at home.

But he puts the emphasis on comfort for patients who are hospitalized because of the rapid progression of pathology, who have a short life expectancy, and who have "no pleasure out of life" or the desire to live.

The fact that parents usually turn first to the antidotal information on labels when a child ingests or inhales a toxic substance is of particular concern to Dr. Rumack. He recommends that anyone facing a poison emergency contact a poison information center before giving any antidote.

Antidotes Listed On Toxic Products Termed Outdated

Medical Tribune Report

DENVER—Many common household products that can be injurious if swallowed or inhaled still carry outdated and possibly harmful information about antidotes on their package labels, a clinical toxicologist warned here.

Dr. Barry H. Rumack, director of the Rocky Mountain Poison Center in Denver General Hospital, cited numerous examples of such faulty antidotal information, including the directives given on containers of Drano, Parson's Ammonia, Easy-Off Oven Cleaner, and Johnson Wax Company's Big Wally.

"The primary problem is that manufacturers of poisons have been improved and upgraded over the years, and many of the companies haven't changed their labels—or at least that portion of their labels—in many years," Dr. Rumack told a conference on critical care held at Swedish Memorial Hospital.

To provide up-to-date information for physicians on potential poisons and their treatment, Dr. Rumack and his colleagues have developed a compendium on more than 100,000 products and compounds. Called "Poisindex," the poison information system is now used at some 125 hospitals and emergency rooms.

In the case of Drano, Dr. Rumack explained that the antidote directive correctly warns against induction of vomiting if the substance has been swallowed. However, the label also advises giving vinegar or a citrus fruit juice. Since Drano is an alkali, the natural acids in these liquids "may cause an increase in the burning of the mouth or esophagus."

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Individualized Therapy Urged in Breast Ca

Medical Tribune Report

HOUSTON—"The availability of many different therapies for a given condition reflect the inadequacy of any single modality," said Dr. Charles K. Tashima Associate Professor of Medicine, The University of Texas Health Science Center at Houston. "Such is the case for breast cancer."

Improvement in treatment can be made by better selecting patients to receive various treatment options: surgery, radiotherapy, endocrine manipulation and chemo-immunotherapy, he told a seminar on the medical management of malignancy sponsored by the M. D. Anderson Tumor Institute.

Dr. Tashima indicated that he favored the Halsted radical mastectomy "when the disease is sufficiently limited in extent so that cure is possible." Less extensive surgical procedures, combined with radiotherapy, represent another alternative. The axilla is not usually irradiated if dissection is adequate, though with significant risk of recurrent disease in the chest wall, the wall will be irradiated, he said.

Dr. Tashima emphasized that treatment should be modified to fit an individual patient, with emphasis in initial presentation on surgery and radiotherapy. Chemo-immunotherapy currently used in his department consists of FAC-BCG (5-FU, Adriamycin and Cytotoxin plus BCG) which produces a response in 75 per cent of patients and median remission of 16 months. Adriamycin is the best single drug in use, Dr. Tashima stated.

He does not recommend prophylactic castration in breast cancer patients nor does he suggest taking women off birth control pills. "Patients with functioning ovaries are probably not affected by small additions of hormones, so that birth control pills are usually not interdicted in menstruating women," he said. However, he does recommend prophylactic castration.

Such a concept would help to determine the recurrence possibility for various groups. He feels it is urgent, since adjuvant chemotherapy significantly affects recurrence rates and survival.

He also feels a method of categorizing patients according to number of sites involved, amount of tumor, and tumor growth rate would further help

to select patients for appropriate therapy, including those patients for whom no treatment is appropriate.

Significant Risk

"In spite of the aggressive approach we have adopted, all the modalities of therapy carry a significant risk and the side effects of treatment are considerable," Dr. Tashima said. For the patient with a small primary and negative axillary nodes, no radiotherapy or adjuvant chemoimmunotherapy is offered, since a radical mastectomy provides a 10-year survival for 80 per cent of patients. He also mentioned the occasional patients with asymptomatic metastatic disease, who have lived with their disease for a number of years. "They appear to have an adequate defense to the tumor and treatment may even be deleterious for these patients."

Though a patient with both primary and metastatic disease may undergo a simple mastectomy, radiotherapy, and chemotherapy, Dr. Tashima again emphasized his belief in the radical procedure, if the choice is simple vs. radical. "There's lots of discussion about things that probably won't make a difference in overall survival. But until more data are in, I would go with the radical," he said.

Lung Disease Mortality Dropped In Fuel Shortage

Medical Tribune Report

BERKELEY, CALIF.—The mortality rate from cardiovascular and chronic lung disease decreased substantially in the San Francisco Bay area during last year's nationwide fuel shortage but returned to normal levels when it ended, a study by Dr. Stephen Brown of the U.C.-Berkeley School of Public Health.

When gasoline sales fell nearly 10 per cent around San Francisco, the death rate from such chronic lung diseases as bronchitis, asthma, and emphysema dropped 33 per cent in San Francisco and 38 per cent in less-urbanized Alameda County. Heart disease deaths also dropped, by 17 and 11 per cent in the respective areas, he said.

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CLINICAL NEWS NOTE: "Physicians have a right to refuse to prescribe the [oral contraceptive] agents if in their best judgment—after reviewing the history, the physical, and the lab tests—they feel the patient's risk is too high. They have to be able to practice according to the dictates of their conscience combined with their best medical judgment." (Dr. Louise B. Tyrer, vice president for medical affairs, Planned Parenthood Federation of America. See page 1.)

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Studies in monkeys of the brain's appetite control centers and the mechanism of satiety are conducted by Dr. Paul R. McHugh at the University of Oregon Health Sciences Center, and have shown that satiety in animals receiving food directly into stomach is influenced more by calories than nature of food.

... brief summaries of editorials or comments in current medical and scientific journals.

The Nondisease Exam

"... If one examines the balance sheet of many physicians ... the cost benefit to the provider in terms of gross and net income from the periodic health examination or screening procedures may be considerable. Many practitioners have too long set the visit fee as the "loss leader," while profits arise from the many laboratory tests appended onto the routine—or not so routine—visit. Similarly, reimbursement formulas as established by hospitals and third parties are frequently such that the hospital could not afford to do fewer laboratory tests; abolishing the admission screening that Korvin questions might well push some already hardpressed institutions further towards the brink!

"With this kind of economic incentive firmly entrenched, one has to be realistic about chances for making patterns of medical care more appropriate. "... we must reevaluate the objectives of the periodic health examination. ... we should spend less time and money searching for what is all too often a nondisease." (Editorial, Thomas L. Deblanco, M.D., and John Noble, M.D., *Ann. Int. Med.* 83:271, Aug., 1975)

Neglected Principle

"Most emergency abdominal operations have a clear primary mission: to save a life. Often the patient is a subpar surgical risk as a direct result of the condition creating the emergency. A self-evident principle should govern the surgeon's behavior in these situations: the life-threatening condition should be corrected by the safest and simplest means. Yet, at times, otherwise level-headed surgeons seem to depart from this 'common' sense.

Example: An eighty year old woman correctly undergoes removal of a gangrenous appendix. At the same operation, the surgeon reduces and repairs a large esophageal hiatus hernia suspected from the findings on the chest x-ray film! The result is an undeservedly uncomplicated postoperative course, from which the surgeon erroneously infers that he did the patient a favor. He ignores the risk, which might have cost the patient's life, of prolonging the operation, operating through an infected field, and correcting a situation unrelated to the emergency and probably present for years without causing symptoms...

"We all have known surgeons, and not all of them young, with the uncanny destructive instinct to do the one additional thing that may lengthen the operation inordinately or lead to postoperative complications and even death. Let us be that surgeon, let us remind ourselves to save life at emergency operations and omit the frills! Who would condone the commercial airline pilot who indulges in aerial acrobatics before safely landing his passenger-filled 747?" (Editorial, Stanley O. Hoerr, M.D., *Amer. J. Surg.* 130:1, July, 1975)

Living better with herself...



Navane® (thiothixene)

PRESCRIBING INFORMATION
Navane® (thiothixene hydrochloride)
Capsules 1 mg, 2 mg, 5 mg, 10 mg, 20 mg
Concentrate 5 mg/ml, Intramuscular 2 mg/ml

Actions: Navane is a psychotropic agent of the thioxanthene series. Navane possesses certain chemical and pharmacological similarities to the piperazine phenothiazine group of drugs, but its mode of action has not been clearly established.

In consideration of the known capability of Navane to precipitate, extreme caution should be exercised in patients with a history of convulsive disorders or those taking a hypnotic agent since a paradoxical increase in the convulsive threshold has been reported with the convulsive threshold although Navane may potentiate the actions of the barbiturates. In the dosage of when Navane is administered concurrently.

Carlson as an adjustment of the dosage of Navane is indicated when Navane is administered with

phenothiazine antihistaminic drugs.

The drug exhibiting rather weak anticholinergic properties, although it usually subsides with continued use, may be given to patients with convulsive disorders, but should be used with caution in patients who are known or suspected to have glaucoma, or who might be exposed to such agents, or to those receiving atropine or related drugs.

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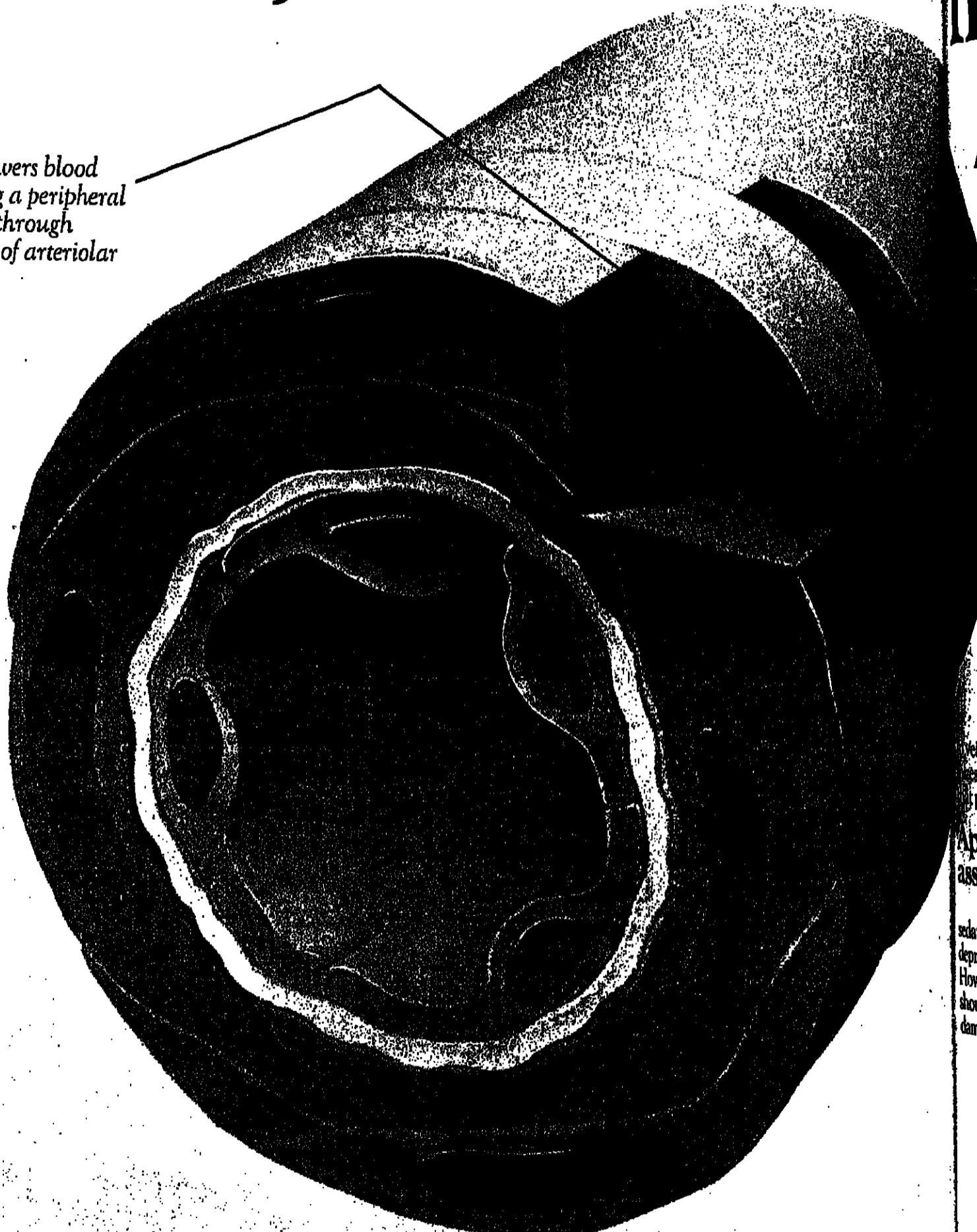
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Apresoline®...where the action is in treating hypertension

Apresoline lowers blood pressure by exerting a peripheral vasodilating effect through a direct relaxation of arteriolar smooth muscle.



An antihypertensive idea whose time has come

Doctors who treat hypertension are increasingly interested in the one oral drug that has a mechanism of action exclusively its own—Apresoline.

Apresoline is in an antihypertensive class by itself because it reduces blood pressure through a unique mechanism. Acting at the ultimate site of hypertension, it directly relaxes arteriolar smooth muscle to decrease peripheral vascular resistance and arterial pressure. As blood pressure falls, there is an accompanying rise in cardiac output and rate.

Apresoline also maintains or increases renal and cerebral blood flow.

Apresoline minimizes postural hypotension

Nickerson¹ describes the action of Apresoline as follows:

"A preferential effect on arterioles, as compared with veins, allows the increase in cardiac output and minimizes postural hypotension; the latter is much less than that produced by agents blocking sympathetic nerves."

Apresoline avoids side effects associated with other agents

Such untoward reactions as drowsiness, lethargy, sedation, sexual dysfunction, and exacerbation of mental depression are not usually encountered with Apresoline. However, as with any antihypertensive agent, hydralazine should be used with caution where advanced renal damage exists.

Apresoline helps tailor the regimen to the patient

When Apresoline is added to an existing antihypertensive regimen, it introduces a different and complementary pharmacologic approach to the control of your patient's hypertension.

Apresoline thus affords the physician a variety of combinations with which he can construct regimens more closely molded to individual requirements. According to Freis,² such a combination of drugs, each with a different antihypertensive mechanism, is the most effective way to control blood pressure. This may also permit lower drug dosages.

Apresoline lends itself admirably to the contemporary antihypertensive rationale and its therapeutic goals: more vigorous and more effective control of blood pressure through a plurality of mechanisms.

Apresoline: used effectively in the VA studies

Apresoline was one of the three basic drugs used in two published VA cooperative studies.^{3,4}

References: 1. Nickerson M: Antihypertensive agents and the drug therapy of hypertension. In Goodman LS, Gilman A (eds): *The Pharmacological Basis of Therapeutics*, 5th ed. New York, The Macmillan Company, 1970, p 729. 2. Freis ED: Hypertension: a cardiovascular disease. *Clin Pharmacol Ther* 13:627-632, 1972. 3. Freis ED: Treatment on morbidity in hypertension: Results in patients with diastolic blood pressures averaging 115 through 129 mm Hg. Veterans Administration Cooperative Study Group on Antihypertensive Agents. *JAMA* 202:1026-1034, 1967. 4. Effects of treatment on morbidity in hypertension. II. Results in patients with diastolic blood pressure averaging 90 through 114 mm Hg. Veterans Administration Cooperative Study Group on Antihypertensive Agents. *JAMA* 213:1143-1152, 1970.

Next page: Apresoline (hydralazine) and the Hypertension Task Force

Apresoline® hydralazine hydrochloride (hydralazine hydrochloride)

TABLETS

Indications: Hypertension, alone or as an adjunct. Hypersensitivity, coronary artery disease, mitral valvular rheumatic heart disease.

Cautions: Administration of doses over 800 mg per day may produce an antidiuretic hydralazine effect.

ing to a clinical picture simulating acute systemic lupus erythematosus. This may also occur at upon withdrawal of therapy, but long-term treatment with steroids may be required.

Hypersensitivity, coronary artery disease, mitral valvular rheumatic heart disease.

Cautions: Administration of doses over 800 mg per day may produce an antidiuretic hydralazine effect.

Usage in Pregnancy:

The drug should be used only when, in the judgment of the physician, it is deemed essential to the welfare of the patient.

Precautions:

Use cautiously in suspected coronary artery or other vascular diseases, cerebral vascular accidents, and atherosclerosis. Hypertension, cerebral vascular, and peripheral nerve damage. Postural hypotension may occur, and the drug may respond to anticholinergics. Discontinuance therapy.

Adverse Reactions:

Common: Headache, tachycardia, anorexia, nausea, vomiting, diarrhea, tachycardia, atrial fibrillation, Less frequent: Nasal congestion, flushed, tachycardia, conjunctivitis, peripheral neuritis.

Induced by hydralazine, numbness, and transient reactions associated with a headache, respiratory, or circulatory reaction.

Induced by hydralazine, consisting of reduction in hemoglobin and red cell count, leukopenia, agranulocytosis, and purpura. These reactions are rare.

Induced by hydralazine, peripheral neuritis.

Tablets, 100 mg (peach, dry-coated); bottles of 100.

Consult complete literature before prescribing.

CIBA Pharmaceutical Company

Division of CIBA Corporation

Summit, New Jersey 07901

C I B A

Apresoline... (hydralazine) part of the Hypertension Task Force "plan of action"

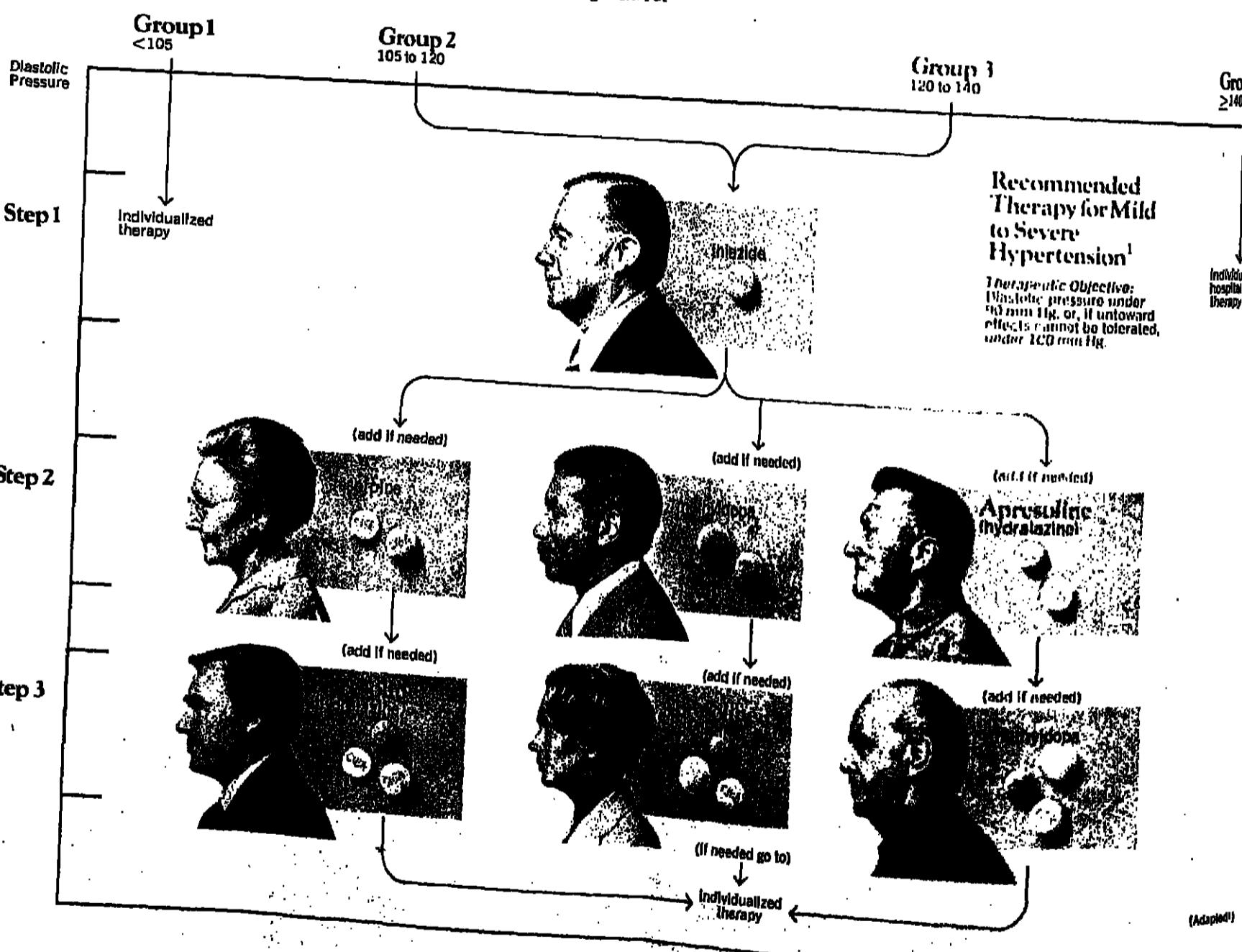
In September 1973, Task Force I of the National High Blood Pressure Education Program recommended a series of antihypertensive regimens for groups with hypertension ranging from mild to severe. Hydralazine—used in combination with sympathetic-inhibiting and/or diuretic antihypertensive

agents—was a specific recommendation for "second step" and "third step" therapy in patients with diastolic pressures ranging from 105 to 140 mm Hg.

Hydralazine played a prominent role in the Task Force regimens because of its compatibility with almost any antihypertensive regimen. For

Apresoline can be combined advantageously with nearly all diuretics and sympathetic inhibitors.

Reference: 1. Report of Task Force I, National High Blood Pressure Education Program. Recommendations for a National High Blood Pressure Program Data Base for Effective Antihypertensive Therapy, Sept 1, 1973, DHEW Publication No. (NIH) 74-593.



Apresoline (hydralazine)
...acts directly at the ultimate
site of hypertension
...brings something
special to almost any
antihypertensive
regimen

For brief prescribing information,
please see preceding pages.

CIBA

Wednesday, November 5, 1975

MEDICAL TRIBUNE

11

The Only Independent Weekly Medical Newspaper in the U.S.

Medical Tribune

and Medical News
Published by Medical Tribune, Inc.

A Watergate-Like Stench

LET US NEVER FORGET that Watergate was not a simple political manipulation but the fundamental violation of American civilian rights—through subversion of the electoral process. Let us not forget that it was carried forward by a regime cloaked in the mantle of "law and order."

And let us now remember that part of the totality of the fraud perpetrated upon the American people was the creation of a drug abuse hysteria.

There is not the slightest doubt that problems of drug abuse exist. But, what was a genuine social ill and medical concern was manipulated for primarily political purposes. The real drug addiction and abuse problems in the United States relate, without any serious contention, to alcohol and cigarettes.

False issues lead to distorted perspectives and as a result a law was passed (Controlled Substances Act of 1970) in which, over the repeated protests of MEDICAL TRIBUNE, important therapeutic agents were stigmatized by association with strong drugs of abuse. Hospitals and doctors became burdened with more red tape paperwork. A new "drug regulatory agency" was created and the control of a huge sector of therapeutics was vested in the Justice Department's Drug Enforcement Agency. MEDICAL TRIBUNE at that time warned of the dangers of a "police" approach to medical problems and the potential threat to the rights of scientists.

Third, as though this were not enough, we are learning that the C.I.A. likewise became involved and in its maneuvers within the Drug Enforcement Agency engaged in a pattern of illegal actions within the country.

Fourth, at a time when it was impossible without forfeiting one's academic rights to continue research on LSD, it is now revealed that a secret program of LSD administration "to unscrupulous subjects to learn its effects" was associated with the death of a high-level CIA researcher in biological warfare. His family is quoted as stating that "Without his knowledge or consent [he had] been given LSD by two C.I.A. employees during [a] research meeting."

Science, and particularly the biomedical sciences, must be protected from abuse as a publicity vehicle or as a political football if we are to continue to have a free and democratic science and a healthier society. A.M.S.

We Must Be Doing Something Right

THE TOP ADMINISTRATORS for health of the Department of Health, Education and Welfare have just issued the second Forward Plan for Health, aimed at the five-year period for fiscal years 1977-1981. We quote from the section on current health status: "After a decade of stable mortality rates in the United States, the age adjusted mortality rates have again shown a steady decline of one per cent per year since

1968. The causes of the leveling off during the preceding decade and the recent renewed downturn are not as yet well understood."

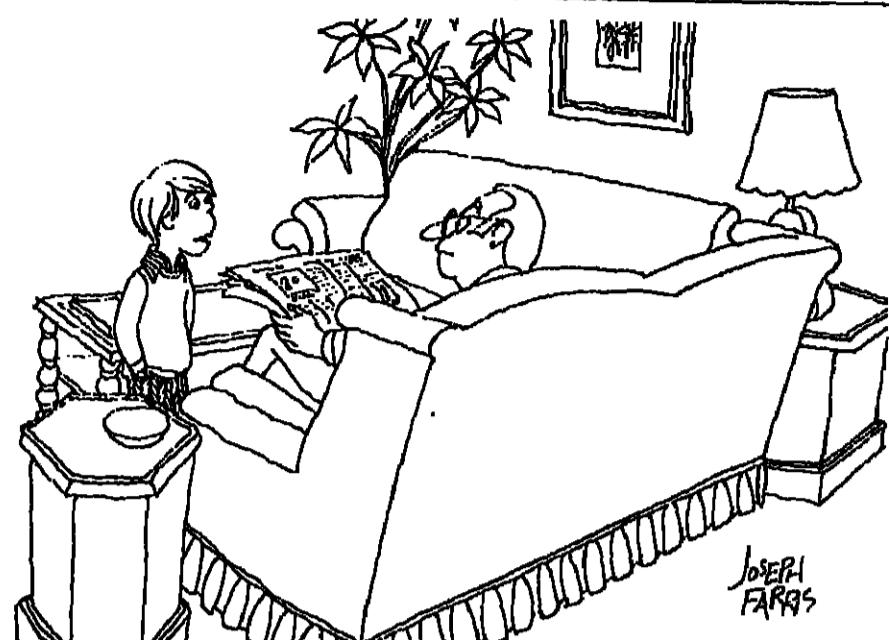
Perhaps the causes "are not as yet

well understood" but as the title of a recent editorial put it, "We must be doing something right. Curiously enough, the critics of health care in this country have not chosen to publicize the good news."

Transposed Great Arteries

CLINICAL QUOTE: "Twenty days after surgery [the infant's] pulmonary pressure was 25/13 mm Hg and the pressure in the right ventricle was 3,700 at the time of surgery." (Dr. A. D. Jatene, Professor of Surgery, Cardiologic Institute, São Paulo, Brazil, describing the first patient ever to undergo total anatomic correction for transposition of the great arteries. See page 1.)

re-evaluated and considered in very good condition, without cyanosis. He presently weighs 5,500 grams (equivalent to 3,700 at the time of surgery). (Dr. A. D. Jatene, Professor of Surgery, Cardiologic Institute, São Paulo, Brazil, describing the first patient ever to undergo total anatomic correction for transposition of the great arteries. See page 1.)



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LETTERS TO TRIBUNE

Shin Splints and Architis

To protect the right
Of the people to know
What the doctors don't know,
Is unsold Sid,
Now making his bid,
For the hero's place—
The consumer's good grace.

Unsold, I say,
Curse I'll bet 'til this day
Sid Wolfe never treated
Diabetes, nor needed
The dastardly pill.

Which he believes can kill;
But the rest of us use
To relieve those who choose
To ask us for relief
Of considerable grief,
Caused by these "polys"
Of untreated diabetes,
And itching of skin
And blurring of vision,
Loss of weight and strength—
I could go on at length.

Far be it from me
To judge U.G.D.P.,
But pills I don't give
To diabetes who live
In a comfortable state
With their glycemic fate,
And do not bemoan
Diet alone.

Could it just be
That U.G.D.P.
May have killed some cases
With Orinase,
By giving the pill
To patients not ill,
Only hyperglycemic—
Then hypoglycemic,
With too much catechol
For a diseased heart's control,
Causing V. Tachy—
Then heart attack.

Sometimes, like the pills,
When insulin kills,
It's hard to see
At the autopsy,
How the patient died
From lack of sucrose.

Oh, we need to design
More waivers to sign!
Nelson G. GOODMAN, M.D.
Bowie, Md.

One to Diabetes Hearings

Your report of the F.D.A. diabetes hearings prompted the following:

Oh, We Need To Design Many
Waivers To Sign

Praise be to Proust
Whom we've heard out
These many years
Arousing fears,
With religious zeal,
Seeking repeat
Of F.D.A. approval,
And near total removal,
Of bad agents
That (?) kill patients.

Now in this crusade,
To save and aid
Dr. Proust in his fight

SPECIFIC SYMPTOM: NONPRODUCTIVE COUGH



SPECIFIC RX: **Hycotuss®** EXPECTORANT

Because specific symptoms require specific therapy, Hycotuss® Expectorant was formulated to specifically treat nonproductive cough associated with respiratory tract congestion.

Hycotuss® Expectorant contains hydrocodone bitartrate, a highly effective antitussive, and glyceryl guaiacolate which acts to liquefy and dislodge viscous secretions in the bronchi.

Relieves persistent coughing while it helps liquefy bronchial secretions.

HYCOTUSS® is a registered U.S. trademark. Where permitted by state laws and regulations.

DESCRIPTION Each 1/2 teaspoonful (6 ml) contains:

Hydrocodone Bitartrate.....5 mg

Glyceryl Guaiacolate.....100 mg

Alcohol U.S.P. 10% v/v

Hydrocodone is 7, 8-dihydrocodeine, a derivative of codeine.

ACTIONS Hydrocodone is a centrally acting narcotic anti-disease providing cough relief for up to 6 hours. Glyceryl guaiacolate exerts its expectorant action by producing a less viscous mucus thereby facilitating its expulsion.

INDICATIONS Indicated for the symptomatic relief of coughs. Especially useful in nonproductive coughs associated with upper and lower respiratory tract congestion.

CONTRAINDICATIONS Hycotuss® Expectorant should not be used in patients with hypersensitivity to hydrocodone or glyceryl guaiacolate.

WARNING Hycotuss® Expectorant should be prescribed and administered with the same degree of caution appropriate for the use of other oral narcotic-containing medications since it is a narcotic. Dependence on it, however, is not a potential for abuse. Patients should be advised not to drive or operate machinery if they become drowsy or show impaired mental and/or physical abilities while taking Hycotuss® Expectorant. Patients receiving narcotic analgesics, phenothiazines, other tranquilizers, sedative-hypnotics or other central nervous system depressants (including alcohol) concomitantly with Hycotuss® Expectorant should be monitored for central nervous system depression. When such combination therapy is employed, the doses of one or both agents should be reduced.

PRECAUTIONS Before prescribing medication to suppress or modify cough, it is important to ascertain that the underlying cause of cough is identified. If medication of bronchitis does not increase the risk of clinical or physiologic bronchitis, and if appropriate therapy for the primary disease is given.

TREATMENT Primary attention should be given to the reestablishment of adequate respiratory exchange through provision of oxygen, if necessary, and maintenance of unobstructed airways. In patients with bronchitis, oxygen, circulatory and respiratory depressions are specific entities requiring appropriate therapy. The bronchitis component requires respiratory depression which may result from oxygen or tissue sensitivity to narcotics, including hydrocodone. An appropriate dose of a new analgesic should be used with caution by the physician, route, administration, and frequency of administration. Since the analgesic component should be kept under close surveillance and never stopped, the practitioner should be advised of the patient's need to monitor hydrocodone requirements.

ADVERSE REACTIONS Adverse reactions, when they occur, include sedation, nausea, vomiting and constipation.

DOSES AND ADMINISTRATION Hycotuss® Expectorant should be taken three times a day and at bedtime, not less than 4 hours apart. Dosages should be altered with the suggested initial dose and subsequent doses adjusted if required.

Usual Dosage:

Adults 1/2 teaspoonful every four hours, after meals and at bedtime.

Children (Over 12 years) same as adults. (2 to 12 years) 1/2 teaspoonful every four hours and at bedtime.

Note: Telephone Rx's may be refilled 5 times within 6 months. Telephone Rx's permitted in most states.

See Brief Summary for prescribing information

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See Brief Summary for prescribing information

Tribune Economic Analysis
Liquidity and Gold Prices
By ELLIOT JANeway
Assistant Professor of Economics
University of Florida College of Medicine

From here on out, the more the rate of inflation jumps, the more it will dry up the liquidity available to everyone.

Roosevelt's purpose in raising the price of gold in the depression of the 1930s was explicitly and unequivocally to undo the ravages of deflation and to start up the corrective momentum of inflation. Roosevelt may have been an economic illiterate—no doubt practical politicians always are. But he grasped the marketplace reality that raising the price of gold primes the pump for inflation—provided liquidity is abundant enough to support the exercise. This was in Roosevelt's time. The opposite is the case now.

Spiraling Interest Rates

The resurgence of inflation is the direct and inescapable reason for the renewed spiraling of interest rates. Today, 7 per cent in tax-exempt income—14 per cent to anyone in a 50 per cent bracket—is no trick for investors willing to tie up money for a year. But money is too scarce and too scared to take advantage of this rate of return. Anytime money is unwilling and unable to accept bonus pay for the privilege of going to work, it's not likely to volunteer for the chance to shoot craps in the gold game.

It's little wonder that the very governments which the gold bugs counted on to pull the price of gold are now breaking it. The liquidity crunch is hurting them most. Governments are under the most urgent and endless pressure to raise cash. The actual announcements of sales from official government holdings are only the tip of the iceberg. No private speculators can hope to support the gold market when distress government selling is breaking it.

Ask Janeway

Would you recommend a retired couple invest the majority of their funds in bonds? We are considering BBB-rated utility bonds and long-term Treasury bonds because income is our highest priority. However, this would use up funds for an extended period of time and would provide only minimum flexibility.

Medical Couple, M.D. & R.N.

What about conserving your capital? Your thinking would expose you to capital losses as interest rates continue to rise. Retirees may regard themselves as realistic in subordinating growth and gain to income, but they are actually being extremely unrealistic. As retirees you will have no chance to earn back any losses this thinking locks you into.

DRUG INTERACTIONS The central nervous system depressants of Hycotuss® Expectorant may be additive with those of other central nervous system depressants (including alcohol) concomitantly with Hycotuss® Expectorant and may also affect the central nervous system depressant effect of Hycotuss® Expectorant. When such combination therapy is employed, the doses of one or both agents should be reduced.

WARNING

MANAGEMENT OF OVERDOSE: Signs and symptoms of serious overdose with Hycotuss® Expectorant may be characterized by respiratory depression, extreme somno-

IN CONSULTATION

What's New and Important in Ophthalmology?

The Consultant

DR. ANTONIO R. GIASSI
Assistant Professor of Ophthalmology
University of Florida College of Medicine

PERHAPS THE GREATEST ADVANCE in the treatment of corneal disease in over a decade has been the development of the soft contact lens and its use as a "bandage." Innumerable cases of severe blinding keratitis (corneal disease) have been cured or controlled through this relatively simple and inexpensive mode of therapy. Patients have been treated who had previously undergone almost all known medical and surgical modalities in an unsuccessful attempt to control the disease process or to restore vision.

In the overwhelming majority of cases, soft lens therapy has been able to provide relief of pain, control of the underlying disease process, to promote the healing of damaged and diseased tissue, and, in many cases, to improve or to restore vision.

vision in patients with corneal ulcers, bullous keratopathy (corneal blisters) or "dry" eyes (a painful condition marked by an insufficiency of tears).

Soft lenses have also been useful in correcting aphakia (the result of cataract surgery). Permanent or constant wear of soft contact lenses in aphakia is not new. Since 1969, we and other ophthalmologists have been using the lutein-cut contact lens for permanent wear in a very selective group of aphakic patients. This procedure has been for cases where its use was absolutely necessary. However, once the patients were selected, most of them were able to wear the lens continuously, 24 hours a day. Not a panacea by any means, it requires a significant amount of knowledge and skill in contact lenses as well as in designing the surgery for this purpose, e.g., round pupil, elimination of corneal astigmatism, etc.

While corneal astigmatism was assumed by some to be the main limiting factor in the wear of soft contact lenses, it has actually been a problem in only about 5 to 10 per cent of the population. More important, in fact crucial, is the fluctuation in vision. These fluctuations are experienced as alternate blurring and clearing of vision. The main cause of vision fluctuation with soft lenses is the fit of the lens. Individual corneas vary in diameter and curvature. Although a soft lens tends to take the shape of the cornea, it may not provide a perfect match even though it may feel comfortable. A poor match with a hard lens would cause corneal swelling, pain and discomfort; with a soft lens it is the vision that suffers.

Hard contact lenses are made of methylmethacrylate or plexiglass, a material that provides excellent visual acuity in a wide range of visual defects. Since soft contact lenses weigh more than hard contact lenses, they have to be fitted slightly larger than the cornea diameter. They range in diameter from 12.3 to 15.5 mm. Hard lenses, in contrast, are presently fitted from 7.00 to 9.00 mm in diameter and don't quite cover the cornea. Due to the fact that these lenses are fitted larger than the cornea, one of the major problems of hard lens wearers—dirt specks or particles lodging under the lens—seldom arises with soft lenses.

While medical use of the bandage lens requires considerable attention and care, it does have advantages.

The main advantage of the procedure stems from the fact that no knife, no sutures, no hospitalization, no nurses, no abnormal financial costs are needed. In addition, if the therapy proves unsuccessful, or if for any reason the physician wishes to terminate therapy, he may do so by the simple procedure of removing the lens from the patient's eye. Moreover, it does not preclude later more risky surgical procedures.

For routine use of contact lenses, when should the hard contact lens be prescribed? When the soft lens?

This should be the choice of the individual, with the proper aid and consultation of his doctor. For optical reasons, there are persons who will see better with hard or soft contact lenses. Since soft lenses take the shape of the cornea, they tend to reproduce corneal astigmatism without correcting it. In these cases, both eye glasses and hard lenses can correct the blurring of vision caused by corneal astigmatism.

Another type of astigmatism, lenticular astigmatism, caused by an irregularity of the natural lens, is corrected only by eye glasses. In many cases, lenticular astigmatism itself corrects corneal astigmatism, eliminating the necessity of correction by accessory lenses. But it is also not uncommon to find that elimination of corneal astigmatism by hard contact lenses will result in the production of residual astigmatism or the bringing out of lenticular astigmatism with its blurring of vision. That this residual astigmatism has not been a significant drawback in the fitting of hard contact lenses stems from the fact that most patients find some degree of residual astigmatism quite tolerable if the other refractive errors are corrected.

After many years and millions of patients wearing contact lenses, both hard and soft lenses have passed the test of time and have proven themselves both safe and effective. Certainly, however, many minor problems still remain with initial and prolonged use of these lenses.

It is said that for every person who successfully adjusts to hard contact lenses, another gives up because of intolerance to the lenses. Although tolerance has been significantly improved with the introduction of the semi-flexible, thin, small, hard contact lens, poor tolerance to a hard foreign body still remains the main problem. In nearly every respect, the soft contact lens is much kinder to the eye than the hard contact lens and is extremely well-tolerated by patients.

Hard contact lenses are more prone to cause corneal edema than soft. When the edema is light and superficial, the patient sees a great cloud over all objects. When the edema becomes more marked, the patient will notice brightly colored halos around light. A poorly fitted hard contact lens can cause a great deal of corneal edema in a relatively short time. Even a well-fitted hard lens can cause corneal edema or injury of corneal epithelium. On the other hand, soft lenses are almost free of this unpleasant side effect. Soft lenses can be worn during all waking hours either from the first day or very shortly after the beginning of adaptation. Hard lens wearers suffer a loss of tolerance for the lens if they don't wear it on a rather regular basis for quite a few hours every day. The soft lens wearer can abandon the lens for as long as he or she wishes and start wearing it again any time without ill effects. Intermittent social wear is another a considerable advantage of this type of lens.

Hard contact lenses, particularly the old, large and thick lenses, when worn for long periods of time can produce a temporary change in the shape of the cornea. Patients are often inconvenienced by this change in vision.

"Let me tell you about the medicine I'm going to prescribe."

TALKING OVER VALIUM®(diazepam) THERAPY WITH YOUR ANXIOUS PATIENT



A patient often benefits by a greater understanding of his treatment program. You may find it helpful to make your patient aware that the purpose of therapy with Valium is to help reduce discomforting and disabling symptoms of excessive psychic tension and anxiety. It is beneficial for him to understand that much of his tension and anxiety can be relieved by your reassurance and counseling, and that these measures can do more than anything else to help him cope with his basic problems. The patient is reassured in knowing he can expect his medication to help him avoid feeling overwhelmed by his symptoms. And it's also good for him to realize that he will be taking Valium only as long as he needs it.

Your expressed confidence in the medication prescribed, and the positive atmosphere in which therapy is given and accepted, work to the patient's advantage.

Selection of a dosage regimen is an important consideration when Valium (diazepam) is prescribed, and dosage should be individualized to achieve maximum beneficial effect. If the patient understands clearly when and how much to take, and if he knows why it's to his benefit to follow the regimen closely, the chances are better that he will take the medication precisely as directed. That should help avoid missed doses and discourage taking too much or too little medication—all of which can have an undesirable effect on the management of the patient's condition.

*"It's important that you
follow my directions
closely."*

*"I'll see you again the week
after next and we'll see
how you're making out."*

portunity to report improvement he has made and to relate whatever continuing or additional difficulties he may be experiencing. It's also a chance for him to describe his response to therapy with Valium.

During follow-up visits, as your patient talks about his medication and about its effects on his symptoms, he will provide the kind of information that will be of great help in evaluating total therapy, adjusting the dosage of Valium, or discontinuing the medication entirely if that seems indicated.

Your patient is often likely to feel reassured when you talk about seeing him again to check his progress. A planned visit evidences your continued interest and affords the patient an opportunity to report improvement he has made and to relate whatever continuing or additional difficulties he may be experiencing. It's also a chance for him to describe his response to therapy with Valium.

Valium®(diazepam)
2-mg, 5-mg, 10-mg scored tablets
for individualized treatment of psychic tension



Please see the following page for a summary of product information.



Valium® (diazepam)

2-mg, 5-mg, 10-mg scored tablets

Prompt, effective action. Valium (diazepam) works rapidly to relieve pronounced psychic tension in patients overreacting to stress and in psychoneurotic patients.

Before prescribing, please consult complete product information, a summary of which follows:

Indications: Tension and anxiety states; somatic complaints which are concomitants of emotional factors; psychoneurotic states manifested by tension, anxiety, apprehension, fatigue, depressive symptoms or agitation; symptomatic relief of acute agitation, tremor, delirium tremens and hallucinosis due to acute alcohol withdrawal; adjunctively in skeletal muscle spasm due to reflex spasm to local pathology; spasticity caused by upper motor neuron disorders; athetosis; stiff-man syndrome; convulsive disorders (not for sole therapy).

Contraindicated: Known hypersensitivity to the drug. Children under 6 months of age. Acute narrow angle glaucoma; may be used in patients with open angle glaucoma who are receiving appropriate therapy.

Warnings: Not of value in psychotic patients.

Caution against hazardous occupations requiring complete mental alertness. When used adjunctively in convulsive disorders, possibility of increase in frequency and/or severity of grand mal seizures may require increased dosage of standard anticonvulsant medication; abrupt withdrawal may be associated with temporary increase in frequency and/or severity of seizures. Advise against simultaneous ingestion of alcohol and other CNS depressants. Withdrawal symptoms (similar to those with barbiturates and alcohol) have occurred following abrupt discontinuance (convulsions, tremor, abdominal and muscle cramps, vomiting and sweating). Keep addiction-prone individuals under careful surveillance because of their predisposition to habituation and dependence. In pregnancy, lactation or women of childbearing age, weigh potential benefit against possible hazard.

Precautions: If combined with other psychotropics or anticonvulsants, consider carefully pharmacology of agents employed; drugs such as phenothiazines, narcotics, barbiturates, MAO inhibitors and other anti-

Wide margin of safety. Valium is generally well tolerated and in usual dosages rarely produces significant adverse reactions. (See prescribing information below.)

Dosage flexibility. Scored Valium 2-, 5-, and 10-mg tablets give you dosage flexibility no tranquilizer capsule can match.

depressants may potentiate its action. Usual precautions indicated in patients severely depressed, or with latent depression, or with suicidal tendencies. Observe usual precautions in impaired renal or hepatic function. Limit dosage to smallest effective amount in elderly and debilitated to preclude ataxia or oversedation.

Side Effects: Drowsiness, confusion, diplopia, hypotension, changes in libido, nausea, fatigue, depression, dysarthria, jaundice, skin rash, ataxia, constipation, headache, incontinence, changes in salivation, slurred speech, tremor, vertigo, urinary retention, blurred vision. Paradoxical reactions such as acute hyperexcited states, anxiety, hallucinations, increased muscle spasticity, insomnia, rage, sleep disturbances, stimulation have been reported; should these occur, discontinue drug. Isolated reports of neutropenia, jaundice; periodic blood counts and liver function tests advisable during long-term therapy.

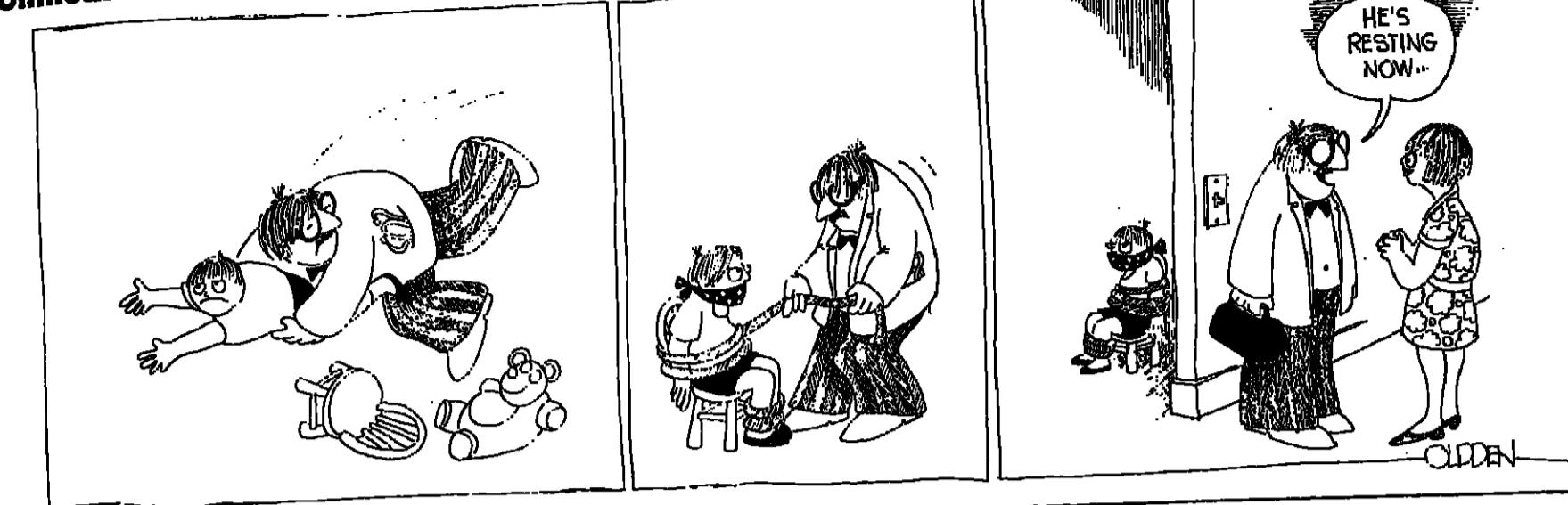
Dosage: Individualize for maximum beneficial effect. **Adults:** Tension, anxiety and psychoneurotic states, 2 to 10 mg b.i.d. to q.i.d.; alcoholism, 10 mg t.i.d. or q.i.d. in first 24 hours, then 5 mg t.i.d. or q.i.d. as needed; adjunctively in skeletal muscle spasm, 2 to 10 mg t.i.d. or q.i.d.; adjunctively in convulsive disorders, 2 to 10 mg b.i.d. to q.i.d. **Geriatric or debilitated patients:** 2 to 2½ mg, 1 or 2 times daily initially, increasing as needed and tolerated. (See Precautions.) **Children:** 1 to 2½ mg t.i.d. or q.i.d. initially, increasing as needed and tolerated (not for use under 6 months).

Supplied: Valium® (diazepam) Tablets, 2 mg, 5 mg and 10 mg—bottles of 100 and 500; Tel-E-Dose® packages of 100, available in trays of 4 reverse-numbered boxes of 25, and in boxes containing 10 strips of 10; Prescription Paks of 50, available singly and in trays of 10.



Roche Laboratories
Division of Hoffmann-La Roche Inc.
Nutley, New Jersey 07110

Clinical Trials



New Combined Drug Held Effective Against All Bacteria Tested

Continued from page 2
bactericidal activity. In fact, the researchers reported, PCS not only eliminates DFA self-reversal but also enhances the antimicrobial activity of both agents manifold.

The new drug has proven equally effective when given to mice orally or by injection against a broad spectrum of bacterial species, including all the serious pathogens for man. The Merck scientists were particularly pleased that *Pseudomonas aeruginosa*, a highly resistant pathogen which is a growing problem in hospitalized patients, proved susceptible to the drug's effect.

According to Dr. Christopher M. Martin, senior director of medical affairs at Merck's research laboratories, not one bacterial strain tested so far has been resistant to the drug. He said the company was "cautiously optimistic that bacteria will have a terrible time with this drug."

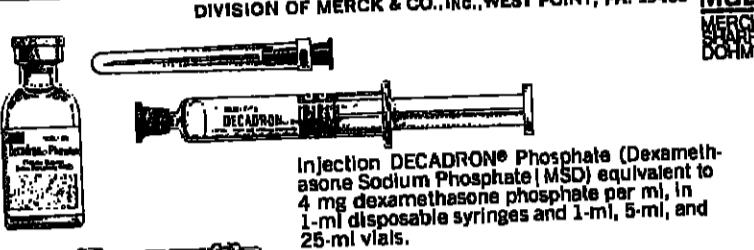
Apprehension that the new agent might kill off harmless and necessary bacteria as well as virulent pathogens has been dispelled by studies in mice which show that it is absorbed into the bloodstream from the upper intestinal tract, Dr. Martin said. Bacteria in the lower tract, the mouth and the skin were unaffected.

Safety testing in human volunteers is expected to begin in early 1976, Dr. Martin announced. Monkeys receiving up to 30 times the normal human dose have exhibited no side effects. However, he cautioned, earlier cycloserine drugs also produced no side effects in animals but caused tremors, behavioral changes and convulsions in humans.

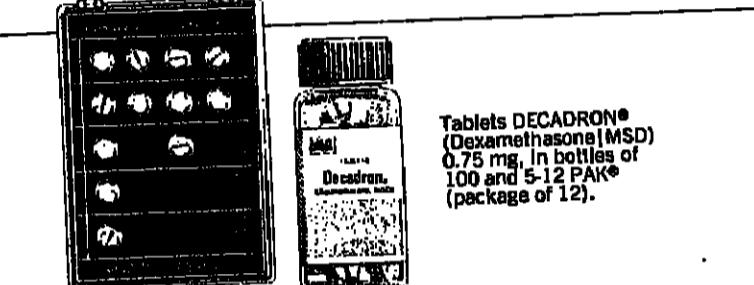
Outpatient Arteriography

Medical Tribune Report
ROCKLAND, MAINE—Outpatient arteriography could mean considerable savings in hospital fees, Drs. Peter E. Giustra and Paul J. Killoran, of the radiology department of Knox County General Hospital, said recently. In a four-year study of 300 patients requiring arteriography, the physicians found no increase in complications and no hospital readmissions among 100 outpatients. The study confirms other reports that complications arise during or right after arteriography.

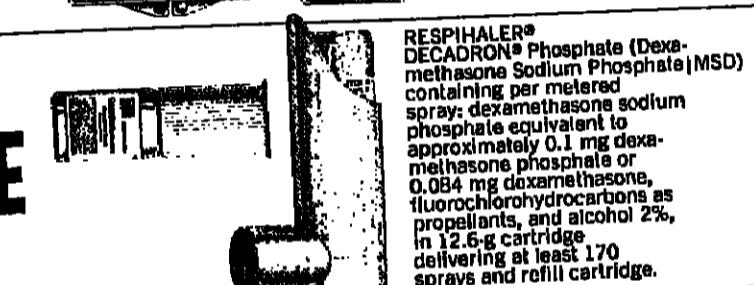
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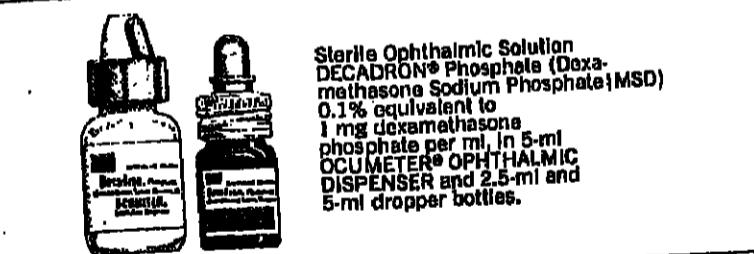
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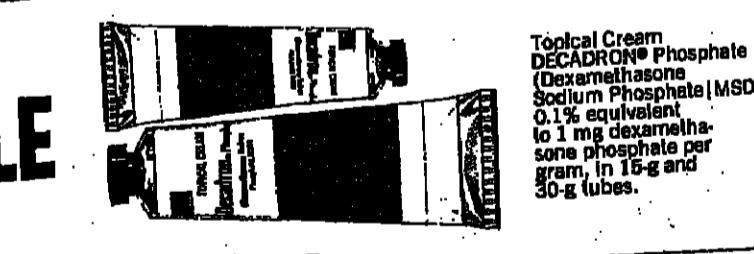
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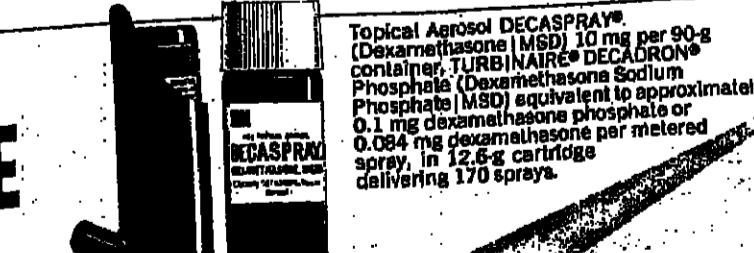
DROPPABLE



SPREADABLE



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Now Suspension
DECADRON-LA®
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equivalent to 8 mg
dexamethasone per ml;
in 5-ml vials.

Major Victory Seen in Grant Policy Reversal

Continued from page 1

O.M.B., ever the dour keeper of the government's purse strings, had vigorously opposed continuation of the main education-funding feature of the 1971-74 law—capitation, under which health schools have been granted specified amounts of money for each student they enroll.

Dr. Mathews, who, as president of the University of Alabama, had worked hand-in-hand with the university's College of Community Health Sciences to resolve health manpower problems in rural Alabama, presumably brought his experience to bear in convincing the White House Domestic Council to endorse the new approach. Dr. Cooper, for his part, was also reportedly dissatisfied with the old line of thinking.

The anticipated legislation, which would replace the 1971-74 health manpower law that expired at the middle of last year, specifies among other things how and to what extent the federal government will finance the training of physicians, osteopaths, dentists, veterinarians, optometrists, pharmacists, podiatrists, and public health specialists.

'Imaginative' and 'Responsive'

Although the first bill to replace the expired statute was introduced a year and a half ago and many others have been put forward since then, no new law has been enacted because of conflicts over ways to support health education, spending levels, and methods of dealing with geographic and specialty maldistribution and the foreign medical graduate (FMG) problem.

Unexpectedly, on September 16 Dr. Cooper outlined a completely new administration proposal.

Senator Kennedy, who with other legislators (including Republicans) had been dismayed by what they called the negativity of previous administration bills, immediately hailed Dr. Cooper's testimony before his Senate health sub-

High Cataract Rate Found In Asthmatic Children On Corticosteroid Drugs

Continued from page 1

"Findings in the children we looked at, when compared to our control of 35 child asthmatics who had not been on oral steroids, are definitely cause for concern," he added.

Although most of the children studied had been taking prednisone, the investigators said they could not incriminate one steroid over another. And despite earlier reports of the possibility that the presence of eczema might have a relationship in cataract formation in patients who are on chronic steroid treatment, the study found no such evidence.

"Clearly, our point is that there may be some children who don't need the doses now being prescribed, and maybe in the future we should look more carefully at the efficacy of day to day steroid therapy," Dr. Spaulding said.

"At any rate," he added, "there is no doubt that children who require corticosteroids to keep their asthma under control should be afforded the opportunity of ophthalmologic examination at least once a year."

committee as "enormously forthcoming," "imaginative," and "responsive."

So effusive were the Senator and Assistant Secretary in their expressed mutual admiration that a subcommittee staff member afterwards called the session "a regular love fest."

Earlier administration bills would have drastically reduced capitation payments as a prelude to abolishing them, but the Senate and House bills that passed their respective chambers last year, but never reached conference, and the refurbished bill that the House passed on July 11 all continued capitation as a basic policy. Now so does the Cooper proposal for schools of medicine, osteopathy, and dentistry, though it reduces capitation payments for student veterinarians, optometrists, and podiatrists and abolishes them for fledgling pharmacists.

H.E.W.'s showdown with O.M.B. and the White House's decision to throw administration support behind capitation occurred only hours before Dr. Cooper was scheduled to testify at the opening session of the Kennedy subcommittee's 1975 hearings on health manpower legislation.

The cliff-hanging nature of the struggle within the administration was such, according to H.E.W. sources, that two prepared testimony statements were actually written for the Assistant Secretary—one favoring capitation and the other opposing it. Some senior H.E.W. officials did not learn which testimony he would present until an hour before he began speaking.

The new administration proposal, which has not yet been formally introduced in either house, would continue capitation at the current level of \$1,500 per year for medical, osteopathic, and dental students, phase it out over the three years of the new law's life (fiscal years 1977 through 1979) if it is enacted by the middle of next year) for veterinary, optometric, and podiatry students, and discontinue it at once for pharmacy students.

It also deals in a much more concrete way than previous administration proposals with the maldistribution and FMG problems that have received so much attention in legislators' bills.

\$55 Million In Scholarships

To remedy the geographic maldistribution of health workers—and particularly doctors—the proposal would require schools to set aside percentages of their enrollments for students who agree to practice in underserved areas after graduation (15 percent in fiscal 1977, 20 percent in fiscal 1978, and 25 percent in fiscal 1979). It would also establish a scholarship program for such students amounting to \$55 million for 5,000 students in fiscal 1977, \$75 million for 7,500 students in fiscal 1978, and \$95 million for 9,500 students in fiscal 1979. Capitation payments to medical, osteopathic, and dental schools that did not agree with these plans would be phased out over the law's life.

The technique emerged in the course of extensive experience with aorto-coronary bypass surgery, Dr. Jatene said, adding: "We believe it is reproducible by most cardiovascular surgeons."

Briefly, in Dr. Jatene's procedure, the ascending aorta and pulmonary trunk are dissected out. The two coronary arteries, along with a piece of the aortic wall, are resected and implanted in the pulmonary artery, still in the posterior

to maintain a specified percentage of their residencies in those specialties (35 per cent in 1977, 40 per cent in 1978, and 50 per cent in 1979).

Though other bills have provisions written into them that would limit the number of FMG's allowed into American residency programs, generally to 25 per cent more than the previous year's total of new graduates from domestic medical schools, the new administration proposal is less specific.

Single Qualifying Exam

"The department...supports the development of a single qualifying examination for all physicians who are entering hospital training programs where they will have some responsibility for patient care," Dr. Cooper said. "We propose to convene appropriate organizations and groups for the development of such an examination. Because this examination is not yet established and because immigrant physicians should be expected to meet standards of U.S. medical graduates in provision of care in the United States, it is proposed that the department will determine the most appropriate screening examination for FMG's."

He also suggested that H.E.W. should develop ways to integrate Americans studying medicine abroad into U.S. medicine through transfer programs.

The new administration measure still differs in some respects from Senator Kennedy's own and that passed by the House in July.

Because the administration, Ken-



Use of new computer-assisted arrhythmia monitor (American Optical) is demonstrated by Dr. Edward A. Partenope and Monica Gelger, R.N., at JFK Medical Center, Edison, N.J.

nedy, and other proposals have still to work their way through the Senate's health subcommittee, the full Labor and Public Welfare Committee, and the Senate itself, it is not considered likely that the Senate will vote out a bill until next spring. When it does, its version of the legislation will have to be reconciled with the House's in conference committee before it goes to the President for signature.

New Procedure Corrects Transposed Great Arteries

Continued from page 1

Jatene stressed that efforts to accomplish total arterial correction of transposed vessels go back to 1954 and 1955 when Drs. Charles C. Bailey and E. B. Kay independently sought to achieve anatomical correction of the defect. Later ingenious attempts to make the correction on the arterial side have been clinical failures, and more recent ideas have not progressed beyond the experimental level, Dr. Jatene noted. In the standard Mustard procedure, the right coronary remains in the pulmonary position. The openings in the aortic wall are closed with a patch. The aorta and pulmonary artery are transected, transposed and then anastomosed. The differences in diameter between the two vessels are equalized by two sutures in the distal and proximal ends of the pulmonary artery so that they correspond to the diameter of the ends of the aorta. The ventricular septal defect is then closed through a right ventriculotomy with a patch.

In Good Condition

In the first clinical trial, the Jatene procedure was performed in an infant with transposed vessels and a large VSD. Hemodynamic studies 20 days after surgery showed complete correction of the defect. The postoperative course was uneventful and the infant at seven months followup is in good condition without cyanosis, Dr. Jatene reported.

Collaborators were Drs. V. G. Fonnes, P. P. Paulista, L. C. B. de Souza, F. Neger, M. Galantier and J. E. M. R. Souza.

In an interview, Dr. Kirklin commented that while there "will be no stampede to use the Jatene technique, it will unquestionably be tried all over the world. Cardiovascular surgeons have been interested for a long time in achieving a correction of this anomaly on the arterial side."

One Man...and Medicine

ARTHUR M. SACKLER, M.D.,
International Publisher, Medical Tribune



To Direct and Not Distract Public Interest Part II

DO NOT THE CONSUMER and so-called "public interest" advocates recognize that unbalanced attacks against doctors and drugs are as dangerous a form of misrepresentation as misleading advertising?

Are they not bound by a higher ethic than that which they feel should apply to the puffery of the propaganda and the actions of vested interests? If the public interest groups represent the interests of the public, should they not lead rather than mislead, should they not direct and not distract?

They cannot have it both ways.

What are the truly major preventable causes of morbidity and mortality in the United States today? Doctors and drugs? The less developed nations of the world do not think so as they suffer the ravages of diseases which are now so rare here they can hardly be demonstrated to medical students in this country.

One would think that implicit in the conclusions of a public interest group calling for cessation of new hospital construction should be the recognition that in an ever-increasingly polluted environment something has happened to reduce morbidity and diminish the need for hospital facilities. Could it be, heaven forbid, doctors and drugs?

They can't have it both ways.

Dangerous Distortions

They are or should be as bound by the ethics of true social responsibility as they must be by the criteria of logic and science. In fact, they should be more sensitive to these than any other group in our society.

They cannot have it both ways.

They are or should be as bound by the ethics of true social responsibility as they must be by the criteria of logic and science. In fact, they should be more sensitive to these than any other group in our society.

Although the patient does all the work, he added, technically the PERC bag is a positive pressure device that works as well as the conventional IPPB.

"This is in essence a positive pressure device, because even the normal breath a person takes is done so under positive atmospheric pressure. Pressure at the mouth is 1,000 centimeters of water, so as the thorax is expanded, thoracic pressure drops to something sub-atmospheric—hence, there exists a

'PERC' Bag Aids Prevention Of Postoperative Atelectasis

Medical Tribune Report

SAN FRANCISCO—An effective, inexpensive device which has proved highly reliable in the prevention of postoperative atelectasis has been developed here by a University of California pulmonary specialist.

Called a perioperative respiratory care (PERC) bag, the device developed by Dr. Anthony Cosentino, director of the pulmonary laboratories at St. Mary's and Mt. Zion Hospitals, and Associate Clinical Professor of Medicine at the University of California, fills the need for a device which allows physicians to calibrate the breaths a postoperative patient takes in efforts to prevent pulmonary complications and gauge necessary amounts of pain medication," Dr. Cosentino told MEDICAL TRIBUNE.

The new device consists of a condom enclosed by a one, one-and-a-half, two or three liter polyethylene bag, which is to be used against significant back pressure, making it effortless for a patient to fill.

According to Dr. Cosentino, the device—which sells to hospitals for about \$2 each—has successfully helped prevent postoperative atelectasis in about 100 patients, without the use of an intermittent positive pressure breathing (IPPB) device.

Patient Does the Work

"We needed an inexpensive bedside device that the patient could operate, and at the same time give us inspiration volume and a good index for amounts of pain medication to administer," Dr. Cosentino said.

Although the patient does all the work, he added, technically the PERC bag is a positive pressure device that works as well as the conventional IPPB.

The technique utilizes a new apparatus which by cooling the surface of the cranium can automatically attain and keep brain temperatures at preselected levels (e.g., 26°C) while moderately cooling the rest of the body to another level (e.g., 30 or 31°C). The apparatus also permits rapid return of temperatures to normal when desired.

Implanted 'Umbrella' Filter Prevents Recurrent Emboli

Continued from page 5

Although the procedure is simple, it is associated with some possible risks and complications, Dr. Schlosser noted. These include errors in placing the filter, shifting of the screen, recurrence of embolism (rarely), and especially edema of the lower extremity.

Dr. Schlosser said patients were generally given constant anticoagulant therapy for one to two years after the operation. To evaluate the hemodynamic situation, 25 patients were examined by angiography over a period of six months to two years.

In eight, the team found complete blocking of the v. cava in the area of the screen filter, with the formation of excessive collaterals, and some tendency to edema of the lower extremity. In eight, the team found complete blocking of the v. cava in the area of the screen filter, with the formation of excessive collaterals, and some tendency to edema of the lower extremity.

Summing it up, the method offers

high measure of protection against pulmonary embolism, with a degree of risk that is tolerable," said Dr. Schlosser.

In certain cases, it could be used prophylactically in phlebographical

confirmed, fresh pelvic and leg vein thromboses, especially in the lower legs, with freely floating thrombi

in the pre-op and post-op phases.

Medicine on Stamps

Joseph Warren



US Bicentennial 10c

Born in Roxbury, Mass., in 1741, he graduated from Harvard in 1759, studied medicine, and very quickly became one of the leading medical men in Boston. Passage of the Stamp Act aroused his patriotic sympathies and he worked diligently in the cause of liberty. Commissioned a major general, he was killed in the battle of Bunker Hill in 1775. He is pictured as the dying soldier in the painting reproduced on the stamp above.

Only the lion and the cock, as Galen says, withstand love's shock. So dearest, do not think me rude if I now yield to lassitude but sympathize with me. I know you would not have me roar, or crow.
Oliver St. John Gogarty, M.D.
(1878-1957)
in After Galen

Stamp: Minkus Publications, Inc., New York
Tint: Dr. Joseph Klein

IN
CONSULTATION

Continued from page 13
enced by their inability to switch back to eye glasses. Changes induced in the cornea by hard lens wear may make the acuity through eye glasses changeable and unsatisfactory for long periods of time after the hard lenses are removed. Experience has shown that complications such as spectacle blur or abrasion occur much less frequently with soft than with hard contact lenses.

Certainly, soft contact lenses are less durable than hard lenses. However, since soft lenses cling to the eye better than the hard ones, they are much less likely to fall out accidentally, not a rare occurrence with hard lenses. Statistics have reported between 25 and 40 per cent of hard contact lens wearers lose one or both of their lenses within the first six months. Of even greater importance is the fact that most of the instances of corneal abrasion, irritation or spectacle blur are produced by a warped, scratched, old, hard contact lens. Both hard and soft contact lenses should be replaced periodically to let the lens wearer benefit from new materials or improvements in technology, and to avoid the damage that can be caused by warped or scratched hard lenses or old soft lenses that have become coated with mucous. It is better not to save the patient's money than to risk potential damage to the eye.

The heat sterilization method employed by one manufacturer of soft lenses and the cold, hydrogen peroxide sterilization procedure employed by another are both extremely safe and effective, with an incidence of clinical bacterial infection certainly not greater than that found in hard contact lenses and perhaps approaching that found in individuals wearing spectacles.

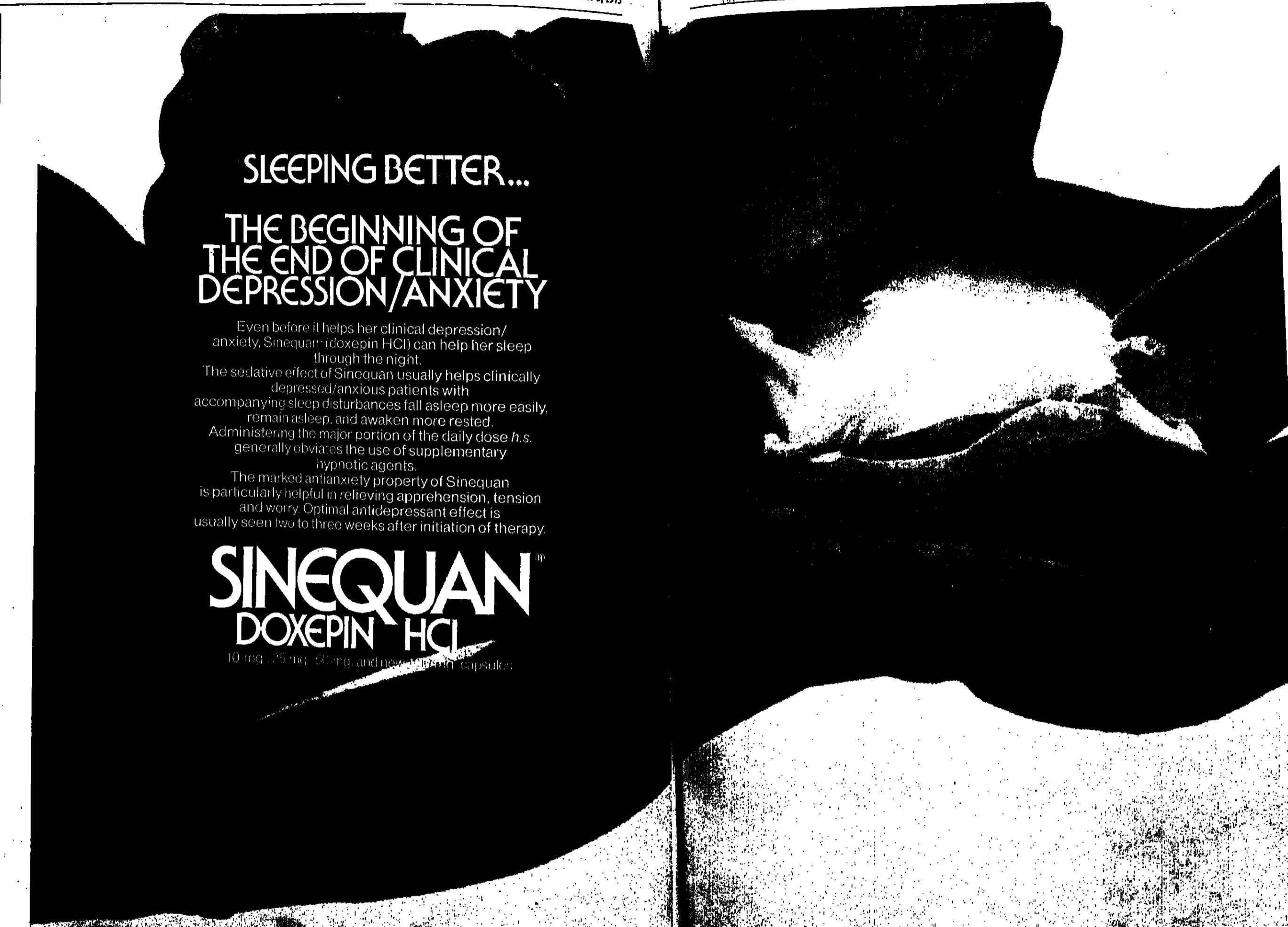
Is there any advantage or disadvantage in having both spectacles and contact lenses?

Eye glasses are the safest, most effective device for the correction of refractive errors. Every contact lens patient should have a pair of spectacles that provide best visual acuity so as to enable free alternation with contact lenses. In addition, when the wearer of hard lenses does put on regular eye glasses and is unable to see clearly, this may be an indication of spectacle blur requiring attention.

A contact lens wearer should be able to switch to eye glasses as necessary for correction and/or comfort, elimination of glare or difficulty seeing at night, or when there are problems of visual acuity, conjunctivitis, irritation, or other complications.

Next In Consultation

DR. JAMES M. STENGLE, Deputy Director for Medical Affairs, Lister Hill National Center for Biomedical Communications, N.I.H., and Chairman, Medical and Scientific Advisory Council, National Hemophilia Foundation, will discuss what's new and important in hemophilia.



SLEEPING BETTER... THE BEGINNING OF THE END OF CLINICAL DEPRESSION/ANXIETY

Even before it helps her clinical depression/anxiety, Sinequan (doxepin HCl) can help her sleep through the night.

The sedative effect of Sinequan usually helps clinically depressed/anxious patients with accompanying sleep disturbances fall asleep more easily, remain asleep, and awaken more rested.

Administering the major portion of the daily dose *h.s.* generally obviates the use of supplementary hypnotic agents.

The marked antianxiety property of Sinequan is particularly helpful in relieving apprehension, tension and worry. Optimal antidepressant effect is usually seen two to three weeks after initiation of therapy.

SINEQUAN DOXEPIH HCl

10 mg, 25 mg, 50 mg, and now 100 mg capsules.

BRIEF SUMMARY Sinequan® (doxepin HCl) Capsules

Contraindications: Sinequan is contraindicated in individuals who have shown hypersensitivity to the drug.

Sinequan is contraindicated in patients with glaucoma or a tendency to urinary retention.

Warnings: Usage in Pregnancy: Sinequan has not been studied in the pregnant patient. It should not be used in pregnant women unless, in the judgment of the physician, it is essential for the welfare of the patient, although animal reproductive studies have not resulted in any teratogenic effects.

Use in Children: The use of Sinequan in children under 12 years of age is not recommended, because safe conditions for its use have not been established.

MAO Inhibitors: Serious side effects and even death have been reported following the concomitant use of certain drugs with MAO inhibitors. Therefore, MAO inhibitors should be discontinued at least two weeks prior to the cautious initiation of therapy with Sinequan (doxepin HCl). The exact length of time may vary and is dependent upon the particular MAO inhibitor being used, the length of time it has been administered, and the dosage involved.

Precautions: Since drowsiness may occur with the use of this drug, patients should be warned of this possibility and cautioned against driving a car or operating dangerous machinery while taking this drug.

Patients: Since alcohol may be potentiated, since alcohol is an inherent risk in any depressed patient and may remain so until

significant improvement has occurred, patients should be closely supervised during the early course of therapy.

Although Sinequan (doxepin HCl) has significant tranquilizing activity, the possibility of activation of psychotic symptoms should be kept in mind.

Other structurally related psychotherapeutic agents (e.g., imidobenzene and dibenzocycloheptenes) are capable of blocking the effects of guanethidine and similarly acting compounds in both the animal and man. Sinequan, however, does not show this effect in animals. At the usual clinical dose, 75 to 150 mg. per day, Sinequan can be given concomitantly with guanethidine and related compounds without blocking the antihypertensive effect. At doses of 300 mg. per day or above, Sinequan does exert a significant blocking effect. In addition,

Sinequan (doxepin HCl) was similar to the other structurally related psychotherapeutic agents as regards its ability to potentiate norepinephrine response in the animal. However, in the human this effect was not seen. This is in agreement with the low incidence of the side effect of tachycardia seen clinically.

Adverse Reactions, Anticholinergic Effects: Dry mouth, blurred vision, and constipation have been reported. They are usually mild, and often subside with continued therapy or reduction of dose.

Central Nervous System Effects: Drowsiness has been observed. This usually occurs early in the course of treatment, and tends to disappear as therapy is continued.

Cardiovascular Effects: Tachycardia and hypertension have been reported infrequently.

Other infrequently reported side effects include extrapyramidal symptoms, gastrointestinal reactions, secretory effects such as increased sweating, weakness, dizziness, fatigue, weight gain, edema, paresthesias, flushing, chills, tinnitus, photophobia, decreased libido, rash, and pruritus.

Dosage: For most patients with illness of mild to moderate severity, a starting dose of 25 mg. t.i.d. is recommended. Dosage may subsequently be increased or decreased at appropriate intervals and according to individual response. The usual optimum dose range is 75 mg./day to 150 mg./day.

In more severely ill patients an initial dose of 50 mg. t.i.d. may be required with subsequent gradual increase to 300 mg./day if necessary. Additional therapeutic effect is rarely to be obtained by exceeding a dose of 300 mg./day.

In patients with very mild symptomatology

IMMATERIA MEDICA

For the President Who Has Nothing

It may be that the President of your favorite medical society, country or club has everything, but just in case you're looking for something, we call your attention to an ad in the Miscellaneous column of the *Wall Street Journal*, sandwiched in between a peat moss ad (200,000 yards) and one for antique hallmarked British silver flatware:

MISCELLANEOUS

FORMER PRESIDENTIAL PRIVATE RAILWAY CAR AVAILABLE Fully equipped, for short or long term lease. Inquiries to be made to Oberleie, Ltd., 280 Madison Avenue, New York City, 10016. (212) 580-1865

The short-term lease idea looked mighty attractive.

Tut-tutted Again

We've been tut-tutted again by Dr. Sam Nixon of Floresville, Texas, because we referred to *The Education of H*Y*M*I*E K*A*P*L*A*N* rather than *H*Y*M*A*N*, which is the way Dr. Sam correctly remembers it.

Our trouble is that we affectionately remember Hyman as Hymie. What can we say? It won't be the first time that affection has led us to err.

But if we may, we'd like to send up a cheer for Leo Rosten who discovered Hymie, Bronx-born and bred, and so hard-up for a tiny bit of recognition that he decorated his name with asterisks. It was probably the most expressive use of typography since e e cummings read archy and mehitabel in *Don Marquis' column*. Let this be considered an "inside" joke accessible only to aging physicians (over 50), we will explain that archy was a cockroach whose physical limitations made it impossible for him to use the typewriter shift key for capitals and punctuation. It was archy who, in archy's new deal, said:

there is bound to be a certain amount of trouble running any country if you are president the trouble happens to you but if you are a tyrant you can arrange things so that most of the trouble happens to other people

